

**Amendments to the Claims:**

This listing of the claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1 (Currently Amended). A method of ~~treating~~  
alleviating the inflammatory response in a subject having an  
inflammatory condition, comprising administering to the  
subject a combination of an anti-inflammatory effective amount  
of methotrexate (MTX) and an anti-inflammatory effective  
amount of an agonist of the A<sub>3</sub> adenosine receptor (A<sub>3</sub>AR  
agonist), wherein the combination provides a combined anti-  
inflammatory effect significantly larger than that provided by  
either MTX or A<sub>3</sub>AR agonist used alone.

2 (Original). A method according to Claim 1,  
wherein MTX is administered to the subject once weekly.

3 (Previously Presented). A method according to  
Claim 1, wherein the A<sub>3</sub>AR agonist is administered to subjects  
between once and a few times a day.

4 (Previously Presented). A method according to  
Claim 1, wherein the A<sub>3</sub>AR agonist is given to the subject  
orally.

5 (Previously Presented). A method according to  
Claim 1, wherein the A<sub>3</sub>AR agonist is IB-MECA or Cl-IB-MECA.

6 (Previously Presented). A method according to

Claim 3, wherein a daily dosage of said A<sub>3</sub>AR agonist is less than 4 mg.

7 (Original). A method according to Claim 6, wherein a daily dosage of said A<sub>3</sub>AR agonist is within the range of about 0.01 to about 2 mg.

8 (Original). A method according to Claim 7, wherein a daily dosage of said A<sub>3</sub>AR agonist is within the range of about 0.1 to about 1.5 mg.

9 (Previously Presented). A method according to Claim 1, wherein the inflammatory condition is an autoimmune disorder.

10 (Currently Amended). A method according to Claim 10, wherein said autoimmune disorder is rheumatoid arthritis.

11. (Original) The method according to Claim 10, wherein said A<sub>3</sub>AR agonist is IB-MECA.

12 (Currently Amended). ~~A~~In a method of treating a subject having an inflammatory condition treatable by methotrexate (MTX), comprising administering to the subject an anti-inflammatory amount of~~and treated with MTX, the improvement comprising administering to the subject an anti-inflammatory effective amount of an agonist of the A<sub>3</sub> adenosine receptor (A<sub>3</sub>AR agonist), wherein the combination of treatment with MTX and A<sub>3</sub>AR agonist provides a combined anti-inflammatory effect significantly larger than that provided by either MTX~~

or A<sub>3</sub>AR agonist used alone.

13 (Currently Amended). ~~A~~In a method of treating a subject having an inflammatory condition treatable by an agonist of the A<sub>3</sub> adenosine receptor (A<sub>3</sub>AR agonist), comprising administering to the subject an anti-inflammatory amount of  
~~and indicated for treatment with an A<sub>3</sub>AR agonist, the~~  
improvement comprising administering to the subject an anti-inflammatory effective amount of methotrexate (MTX), wherein the combination of treatment with MTX and A<sub>3</sub>AR agonist provides a combined anti-inflammatory effect significantly larger than that provided by either MTX or A<sub>3</sub>AR agonist used alone.

14 (Previously Presented). A method according to Claim 12, wherein the A<sub>3</sub>AR agonist is administered to the subject between once and a few times a day.

15 (Previously Presented). A method according to Claim 12, wherein the A<sub>3</sub>AR agonist is given to the subject orally.

16 (Previously Presented). A method according to Claim 12, wherein the A<sub>3</sub>AR agonist is IB-MECA or Cl-IB-MECA.

17 (Previously Presented). A method according to Claim 12, wherein a daily dosage of said A<sub>3</sub>AR agonist is less than 4 mg.

18 (Original). A method according to Claim 17, wherein a daily dosage of said A<sub>3</sub>AR agonist is within the range

of about 0.01 to about 2 mg.

19 (Original). A method according to Claim 18, wherein a daily dosage of said A<sub>3</sub>AR agonist is within the range of about 0.1 to about 1.5 mg.

20 (Previously Presented). A method according to Claim 12, wherein the inflammatory condition is an autoimmune disorder.

21 (Original). A method according to Claim 20, wherein said autoimmune disorder is rheumatoid arthritis.

Claims 22-36 (Cancelled).

37 (New). A method in accordance with claim 1, wherein said anti-inflammatory condition is one in which treatment with MTX is indicated.

38 (New). A method in accordance with claim 37, wherein said anti-inflammatory condition is selected from the group consisting of psoriasis, psoriatic arthritis, Crohn's disease, rheumatoid arthritis, polymyositis and systemic lupus erythematosus.